

AMENDMENTS TO THE CLAIMS:

This listing of claims will replace all prior versions, and listings, of claims in the application:

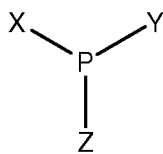
Listing of Claims:

1. (Currently Amended) A chiral transition metal catalyst comprising at least two ~~structurally~~ constitutionally different monophosphorus ligands bonded to a transition metal, at least one monophosphorus ligand being chiral.

2. (Withdrawn) A catalyst as claimed in claim 1, wherein precisely one monophosphorus ligand is chiral.

3. (Original) A catalyst as claimed in claim 1, wherein at least two monophosphorus ligands are chiral.

4. (Previously Presented) A catalyst as claimed in claim 1, wherein the monophosphorus ligands are each independently of the A type



A

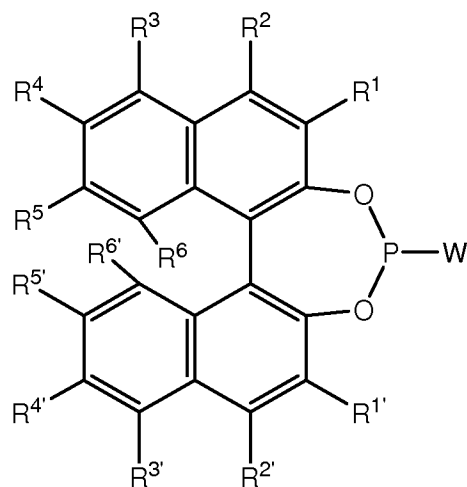
where the X, Y and Z atoms are each independently from the group of carbon, nitrogen, oxygen, sulfur or halogen, to which, according to their number of free valences, further atoms or groups of atoms are bonded independently of one another,

where X, Y and Z may also be connected to one another by the bonded atoms or groups of atoms,

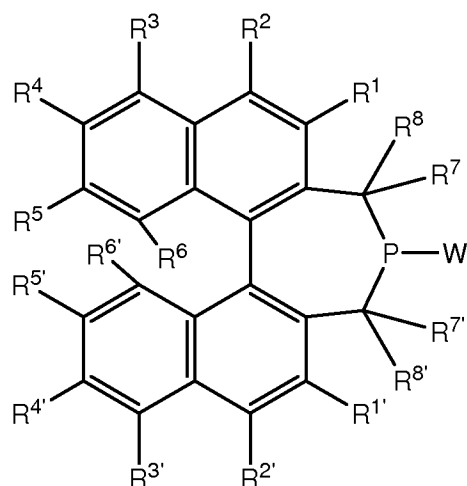
where X-P-Y may also be part of an aromatic system, in which case X is bonded to P by a double bond and there is no substituent Z.

5. (Previously Presented) A catalyst as claimed in claim 1, wherein the monophosphorus ligands are phosphines, phosphites, phosphonites, phosphinites, phosphorous triamides, phosphorous monoester diamides, phosphorous diester amides, phosphonous diamides, phosphinous amides, phosphonous monoester amides, phosphorous halides, phosphorous diamide halides, thiophosphites, thiophosphorous triesters, thiophosphorous monoester diamides or thiophosphorous diesteramides.

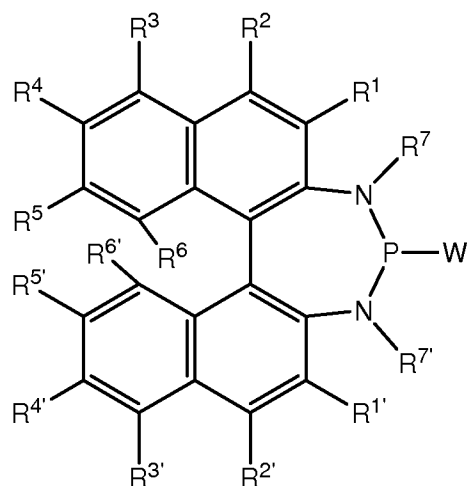
6. (Previously Presented) A catalyst as claimed in claim 1, wherein the chiral ligands are monophosphorus compounds of the B, C or D type



B



C



D

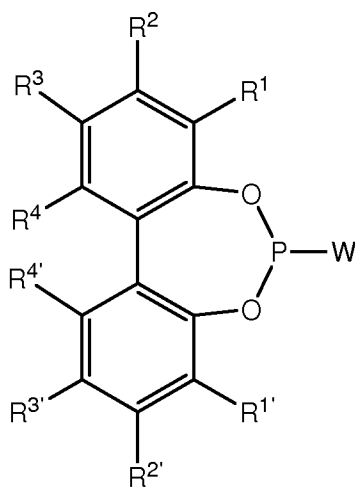
where W is carbon, nitrogen, oxygen, sulfur or halogen, and further atoms or groups of atoms are bonded to W according to its number of free valences,

and where the R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , $R^{1'}$, $R^{2'}$, $R^{3'}$, $R^{4'}$, $R^{5'}$, $R^{6'}$, $R^{7'}$ and $R^{8'}$ radicals are each

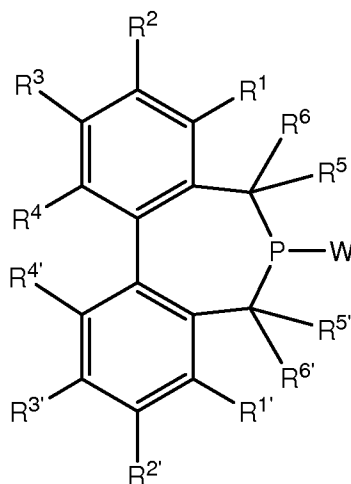
independently from the group of hydrogen, halogen, saturated and unsaturated, linear and branched C₁-C₅₀ alkyl, C₁-C₅₀ aryl, C₁-C₅₀ heteroaryl, alkynyl, silyl, nitro, nitrile, ester, carboxyl, carbonyl, amide, amine, hydroxyl, alkoxy, sulfide and selenide groups, where R¹, R², R³, R⁴, R⁵, R⁶, R⁷, R⁸, R^{1'}, R^{2'}, R^{3'}, R^{4'}, R^{5'}, R^{6'}, R^{7'} and R^{8'} in turn bear further substituents or may be functionalized,

and where one or more carbon atoms of the binaphthyl skeletons may each independently be replaced by the heteroatoms Si, O, N or S.

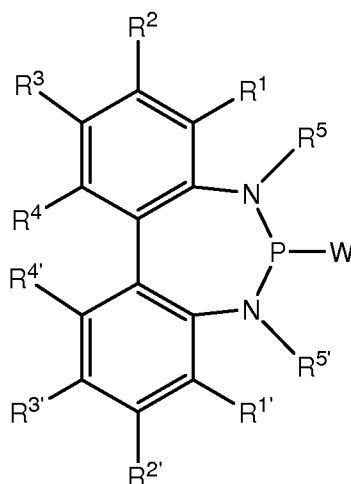
7. (Withdrawn) A catalyst as claimed in claim 1, wherein the chiral ligands are monophosphorus compounds of the E, F or G type



E



F



G

where W is carbon, nitrogen, oxygen, sulfur or halogen, and further atoms or groups of atoms are bonded to W according to its number of free valences,

and where the R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , $R^{1'}$, $R^{2'}$, $R^{3'}$, $R^{4'}$, $R^{5'}$ and $R^{6'}$ radicals are each independently from the group of hydrogen, halogen, saturated and unsaturated, linear and branched C_1 - C_{50}

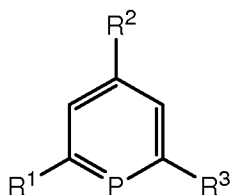
alkyl, C₁-C₅₀ aryl, C₁-C₅₀ heteroaryl, alkynyl, silyl, nitro, nitrile, ester, carboxyl, carbonyl, amide, amine, hydroxyl, alkoxy, sulfide and selenide groups,

where R¹, R², R³, R⁴, R⁵, R⁶, R^{1'}, R^{2'}, R^{3'}, R^{4'}, R^{5'} and R^{6'} in turn bear further substituents or may be functionalized,

and where one or more carbon atoms of the biphenyl skeletons may each independently be replaced by the heteroatoms Si, O, N or S.

8. (Withdrawn) A catalyst as claimed in claim 1, wherein at least one achiral ligand is a monophosphorus compound of the H-T type

PR^1_2R^2	$\text{P}(\text{OR}^1)_2(\text{OR}^2)$	$\text{P}(\text{NR}^1\text{R}^2)_3$
H	I	J
$\text{P}(\text{SR}^1)_2(\text{SR}^2)$	$\text{R}^1\text{OP}(\text{NR}^2\text{R}^3)_2$	$(\text{R}^1\text{O})_2\text{P}(\text{NR}^2\text{R}^3)$
K	L	M
$\text{O}=\text{P}(\text{OR}^1)_2(\text{OR}^2)$	$\text{O}=\text{P}(\text{NR}^1\text{R}^2)_2(\text{NR}^3\text{R}^4)$	$\text{S}=\text{P}(\text{OR}^1)_2(\text{OR}^2)$
N	O	P
$\text{S}=\text{P}(\text{NR}^1\text{R}^2)_2(\text{NR}^3\text{R}^4)$	$\text{R}-\text{N}=\text{P}(\text{OR}^1)_2(\text{OR}^2)$	$\text{R}-\text{N}=\text{P}(\text{NR}^1\text{R}^2)_2(\text{NR}^3\text{R}^4)$
Q	R	S



T

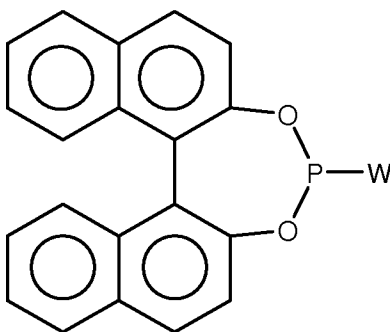
where the R^1 , R^2 , R^3 and R^4 radicals are each independently from the group of hydrogen, halogen, saturated and unsaturated, linear and branched C_1 - C_{50} alkyl, C_1 - C_{50} aryl, C_1 - C_{50} heteroaryl, alkynyl, silyl, nitro, nitrile, ester, carboxyl, carbonyl, amide and selenide groups,

where R^1 , R^2 , R^3 and R^4 in turn bear further substituents, and may be functionalized or bridged.

9. (Previously Presented) A catalyst as claimed in claim 1, wherein the transition metal is a metal of groups IIIb, IVb, Vb, VIb, VIIb, VIII, Ib or IIb of the periodic table or a lanthanide or actinide.

10. (Original) A catalyst as claimed in claim 9, wherein the transition metal is Rh, Ir, Ru, Ni, Pd or Pt.

11. (Currently Amended) A catalyst as claimed in claim 3, wherein the chiral monophosphorus ligands used are at least two ligands of the



type where W is each independently CH_3 , $\underline{C}(CH_3)_3$, $C-C_6H_{11}$ or OCH_3 .

12. (Withdrawn) A process for catalytically preparing chiral organic compounds, comprising conducting a chemical reaction of prochiral organic compounds in the presence of a transition metal catalyst as claimed in claim 1.

13. (Withdrawn) The process as claimed in claim 12, wherein the chemical reaction is a hydrogenation.

14. (Withdrawn) The process as claimed in claim 12, wherein the chemical reaction is a hydroformylation.

15. (Withdrawn) The process as claimed in claim 12, wherein the chemical reaction is a hydroboration, hydrosilylation, hydrovinylation, hydroamination, epoxidation, hydroxylation, aminohydroxylations, substitution, allyl substitution, Heck coupling, Stille coupling, Suzuki coupling, Negishi coupling, Michael addition, aldol addition, Diels-Alder reaction, cyclopropanation, CH insertion reaction or 1,3-dipolar cycloaddition.